

# **EXHIBIT 91**

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# Supplemental Guidance to RAGS: Calculating the Concentration Term

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The overarching mandate of the Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA) is to protect human health and the environment from current and potential threats posed by uncontrolled releases of hazardous substances. To help meet this mandate, the U.S. Environmental Protection Agency's (EPA's) Office of Emergency and Remedial Response has developed a human health risk assessment process as part of its remedial response program. This process is described in the *Risk Assessment Guidance for Superfund: Volume I — Human Health Evaluation Manual (RAGS/HHEM)*. Part A of RAGS/HHEM addresses the baseline risk assessment, and describes a general approach for estimating exposure to individuals from hazardous substance releases at Superfund sites.

This bulletin explains the concentration term in the exposure/intake equation to remedial project managers (RPMs), risk assessors, statisticians, and other personnel. This bulletin presents the general intake equation as presented in RAGS/HHEM Part A, discusses basic concepts concerning the concentration term, describes generally how to calculate the concentration term, presents examples to illustrate several important points, and lastly, identifies where to get additional help.

## THE CONCENTRATION TERM

### How is the concentration term used?

RAGS/HHEM Part A presents the Superfund risk assessment in four "steps": (1) data collection and evaluation; (2) exposure assessment; (3) toxicity assessment; and, (4) risk characterization. The concentration term is calculated for use in the exposure assessment step. **Highlight 1** presents the general equation Superfund uses for calculating exposure, and illustrates that the concentration term (C) is one of several parameters needed to estimate contaminant intake for an individual.

For Superfund assessments, the concentration term (C) in the intake equation is an estimate of the arithmetic average concentration for a contaminant based on a set of site sampling results. Because of the uncertainty associated with estimating the true average concentration at a site, the 95 percent upper confidence limit (UCL) of the arithmetic mean should be used for this variable. The 95 percent UCL provides reasonable confidence that the true site average will not be underestimated.

### Why use an average value for the concentration term?

An estimate of average concentration is used because:

*Supplemental Guidance to RAGS* is a bulletin series on risk assessment of Superfund sites. These bulletins serve as supplements to *Risk Assessment Guidance for Superfund: Volume I—Human Health Evaluation Manual*. The information presented is intended as guidance to EPA and other government employees. It does not constitute rulemaking by the Agency, and may not be relied on to create a substantive or procedural right enforceable by any other person. The Government may take action that is at variance with these bulletins.

**Highlight 1**  
**GENERAL EQUATION FOR ESTIMATING EXPOSURE**  
**TO A SITE CONTAMINANT**

$$I = C \times \frac{CR \times EFD}{BW} \times \frac{1}{AT}$$

where:

I	=	Intake (i.e., the quantitative measure of exposure in RAGS/HHEM)
C	=	Contaminant Concentration
CR	=	Contact (Intake) Rate
EFD	=	Exposure Frequency and Duration
BW	=	Body Weight
AT	=	Averaging Time

- (1) carcinogenic and chronic noncarcinogenic toxicity criteria<sup>1</sup> are based on lifetime average exposures; and,
- (2) Average concentration is most representative of the concentration that would be contacted at a site, over time.

For example, if you assume that an exposed individual moves randomly across an exposure area, then the spatially-averaged soil concentration can be used to estimate the true average concentration contacted over time. In this example, the average concentration contacted over time would equal the spatially averaged concentration over the exposure area. While an individual may not actually exhibit a truly random pattern of movement across an exposure area, the assumption of equal time spent in different parts of the area is a simple but reasonable approach.

**When should an average concentration be used?**

The two types of exposure estimates now being required for Superfund risk assessments, a reasonable maximum exposure (RME) and an average, should both use an average concentration. To be protective, the overall estimate of intake (see **Highlight 1**) used as a basis for action at

Superfund sites should be an estimate in the high-end of the intake/dose distribution. One high-end option is the RME used in the superfund program. The RME, which is defined as the highest exposure that could reasonably be expected to occur for a given exposure pathway at a site, is intended to account for both uncertainty in the contaminant concentration and variability in exposure parameters (e.g., exposure frequency, averaging time). For comparative purposes, agency guidance (U.S. EPA, *Guidance on Risk Characterization for Risk Managers and Risk Assessors*, February 26, 1992) states that an average estimate of exposure also should be presented in risk assessments. For decision-making purposes in the Superfund program, however, RME is used to estimate risk.<sup>2</sup>

**Why use an estimate of the arithmetic mean rather than the geometric mean?**

The choice of the arithmetic mean concentration as the appropriate measure for estimating exposure derives from the need to estimate an individual's long-term average exposure. Most Agency health criteria are based on the long-term average daily dose, which is simply the sum of all daily doses divided by the total number of days in the averaging period. This is the definition of an arithmetic mean. The

<sup>1</sup> When acute toxicity is of most concern, a long-term average concentration generally should not be used for risk assessment purposes, as the focus should be to estimate short-term, peak concentrations.

<sup>2</sup> For additional information on RME, see RAGS/HHEM Part A and the National Oil and Hazardous Substances Pollution contingency plan (NCP), 55 *Federal Register* 8710, March 8, 1990.

arithmetic mean is appropriate regardless of the pattern of daily exposures over time, or the type of statistical distribution that might best describe the sampling data. The geometric mean of a set of sampling results, however, bears no logical connection to the cumulative intake that would result from long-term contact with the site contaminants, and it may differ appreciably from—and be much lower than—the arithmetic mean. Although the geometric mean is a convenient parameter for describing central tendencies of lognormal distributions, it is not an appropriate basis for estimating the concentration term used in Superfund exposure assessments. The following simple example may help clarify the difference between the arithmetic and geometric mean, when used for an exposure assessment:

Assume the daily exposure for a trespasser subject to random exposure at a site is 1.0, 0.01, 1.0, 0.01, 1.0, 0.01, 1.0, and 0.01 units/day, over an 8-day period. Given these values, the cumulative exposure is simply their summation, or 4.04 units. Dividing this by 8 days of exposure results in an arithmetic mean of 0.505 units per day. This is the value we would want to use in a risk assessment for this individual, not the geometric mean of 0.1 units per day. Viewed another way, multiplication of the geometric mean by the number of days equals 0.8 units, considerably lower than the known cumulative exposure of 4.04 units.

## UCL AS AN ESTIMATE OF THE AVERAGE CONCENTRATION

### What is a 95 percent UCL?

The 95 percent UCL of a mean is defined as a value that, when calculated repeatedly for randomly drawn subsets of site data, equals or exceeds the true mean 95 percent of the time. Although the 95 percent UCL of the mean provides a conservative estimate of the average (or mean) concentration, it should not be confused with a 95<sup>th</sup> percentile of site concentration data (as shown in **Highlight 2**).

### Why use the UCL as the average concentration?

Statistical confidence limits are the classical tool for addressing uncertainties of a distribution average. The 95 percent UCL of the arithmetic

mean concentration is used as the average concentration, because it is not possible to know the true mean. The 95 percent UCL, therefore, accounts for uncertainties due to limited sampling data at Superfund sites. As sampling data become less limited at a site, uncertainties decrease, the UCL moves closer to the true mean, and exposure evaluations using either the mean or the UCL produce similar results. This concept is illustrated in **Highlight 2**.

### Should a value other than the 95 percent UCL be used for the concentration?

A value other than the 95 percent UCL can be used, provided the risk assessor can document that high coverage of the true population mean occurs (i.e., the value equals or exceeds the true population mean with high probability). For exposure areas with limited amounts of data or extreme variability in measured or modeled data, the UCL can be greater than the highest measured or modeled concentration. In these cases, if additional data cannot practicably be obtained, the highest measured or modeled value could be used as the concentration term. Note, however, that the true mean still may be higher than this maximum value (i.e., the 95 percent UCL indicates a higher mean is possible), especially if the most contaminated portion of the site has not been sampled.

## CALCULATING THE UCL

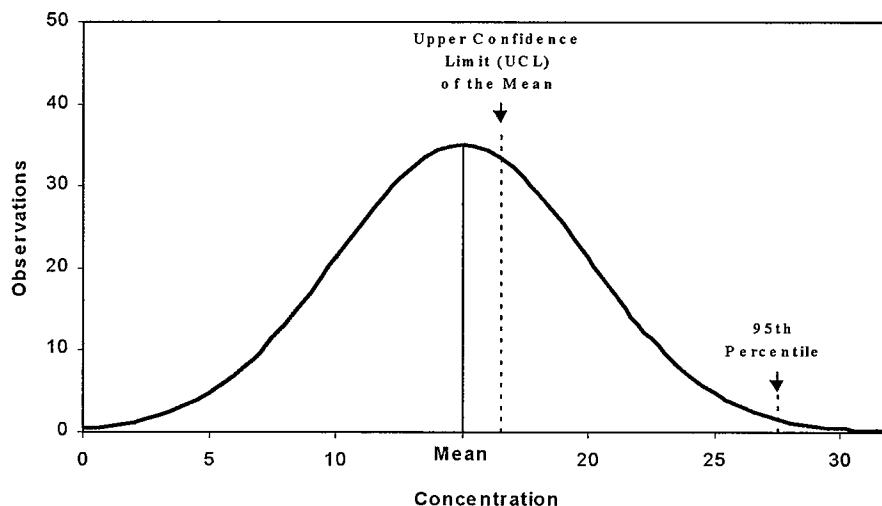
### How many samples are necessary to calculate the 95 percent UCL?

Sampling data from Superfund sites have shown that data sets with fewer than 10 samples per exposure area provide poor estimates of the mean concentration (i.e., there is a large difference between the sample mean and the 95 percent UCL), while data sets with 10 to 20 samples per exposure area provide somewhat better estimates of the mean, and data sets with 20 to 30 samples provide fairly consistent estimates of the mean (i.e., the 95 percent UCL is close to the sample mean). Remember that, in general, the UCL approaches the true mean as more samples are included in the calculation.

### Should the data be transformed?

EPA's experience shows that most large or "complete" environmental contaminant data sets

**Highlight 2**  
**COMPARISON OF UCL AND 95<sup>th</sup> PERCENTILE**



As sample size increases, the UCL of the mean moves closer to the true mean, while the 95<sup>th</sup> percentile of the distribution remains at the upper end of the distribution.

from soil sampling are lognormally distributed, rather than normally distributed (see **Highlights 3 and 4**, for illustrations of lognormal and normal distributions). In most cases, it is reasonable to assume that Superfund soil sampling data are lognormally distributed. Because transformation is a necessary step in calculating the UCL of the arithmetic mean for a lognormal distribution, the data should be transformed by using the natural logarithm function (i.e., calculate  $\ln(x)$ , where  $x$  is the value from the data set). However, in cases where there is a question about the distribution of the data set, a statistical test should be used to identify the best distributional assumption for the data set. The W-test (Gilbert, 1987) is one statistical method that can be used to determine if a data set is consistent with a normal or lognormal distribution. In all cases, it is valuable to plot the data to better understand the contaminant distribution at the site.

**How do you calculate the UCL for a lognormal distribution?**

To calculate the 95 percent UCL of the arithmetic mean for a lognormally-distributed data

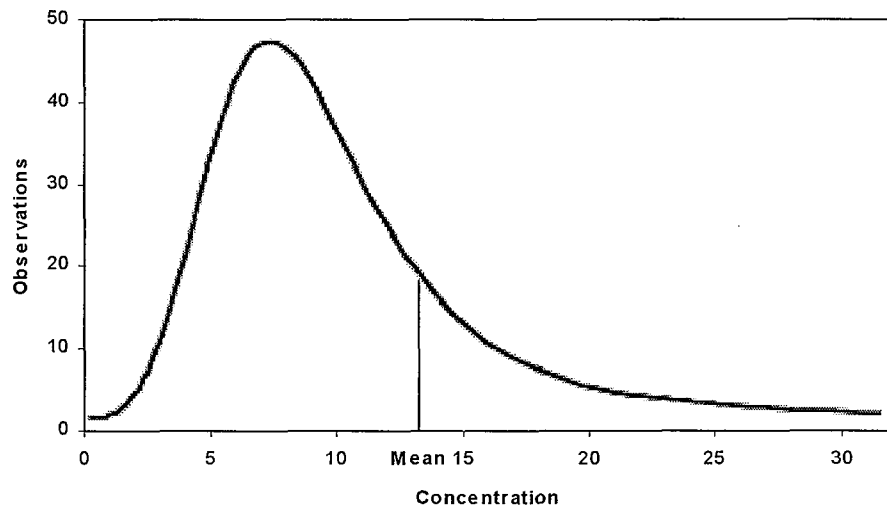
set, first transform the data using the natural logarithm function as discussed previously (i.e., calculate  $\ln(x)$ ). After transforming the data, determine the 95 percent UCL for the data set by completing the following four steps:

- (1) Calculate the arithmetic mean of the transformed data (which is also the log of the geometric mean);
- (2) Calculate the standard deviation of the transformed data;
- (3) Determine the H-statistic (e.g., see Gilbert, 1987); and,
- (4) Calculate the UCL using the equation shown in **Highlight 5**.

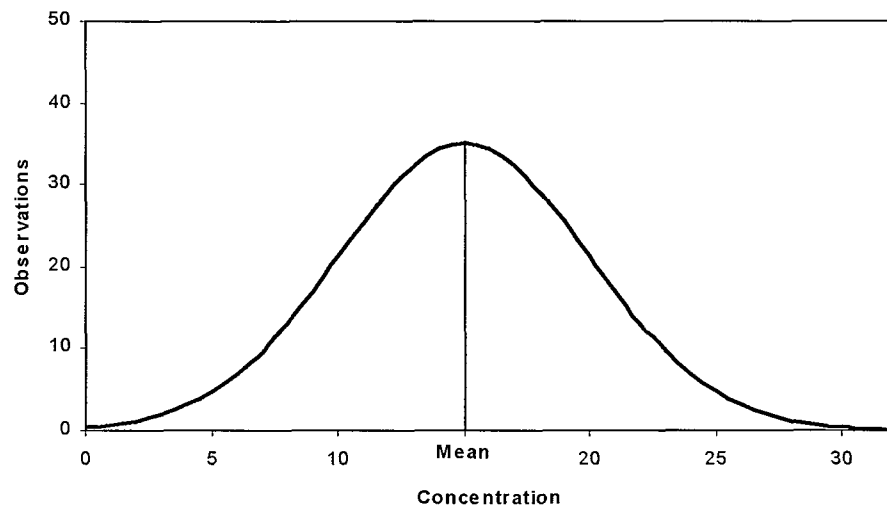
**How do you calculate the UCL for a normal distribution?**

If a statistical test supports the assumption that the data set is normally distributed, calculate the 95 percent UCL by completing the following four steps:

**Highlight 3**  
**EXAMPLE OF A LOGNORMAL DISTRIBUTION**



**Highlight 4**  
**EXAMPLE OF A NORMAL DISTRIBUTION**



**Highlight 5**  
**CALCULATING THE UCL OF THE ARITHMETIC MEAN**  
**FOR A LOGNORMAL DISTRIBUTION**

$$UCL = e^{\left(\bar{x} + 0.5s^2 + sH / \sqrt{n-1}\right)}$$

where:

UCL	=	upper confidence limit
e	=	constant (base of the natural log, equal to 2.718)
$\bar{x}$	=	mean of the transformed data
s	=	standard deviation of the transformed data
H	=	H-Statistic (e.g., from table published in Gilbert, 1987)
n	=	number of samples

**Highlight 6**  
**CALCULATING THE UCL OF THE ARITHMETIC MEAN FOR A NORMAL DISTRIBUTION**

$$UCL = \bar{x} + t(s / \sqrt{n})$$

where:

UCL	=	upper confidence limit
$\bar{x}$	=	mean of the untransformed data
s	=	standard deviation of the untransformed data
t	=	Student-t statistic (e.g., from table published in Gilbert, 1987)
n	=	number of samples

- (1) Calculate the arithmetic mean of the untransformed data;
- (2) Calculate the standard deviation of the untransformed data;
- (3) Determine the one-tailed t-statistic (e.g., see Gilbert, 1987); and,
- (4) Calculate the UCL using the equation shown in **Highlight 6**.

Use caution when applying normal distribution calculations, if there is a possibility that heavily contaminated portions of the site have not been adequately sampled. In such cases, a UCL from normal distribution calculations could fall below the true mean, even if a limited data set at a site appears normally distributed.

## EXAMPLES

The examples show in **Highlights 7 and 8** address the exposure scenario where an individual at a Superfund site has equal opportunity to contact soil in any sector of the contaminated area over time. Even though the examples address only soil exposures, the UCL approach is applicable to all exposure pathways. Guidance and examples for other exposure pathways will be presented in forthcoming bulletins.

**Highlight 7** presents a simple data set and provides a stepwise demonstration of transforming the data—assuming a lognormal distribution—and calculating the UCL. **Highlight 8** uses the same data set to show the difference between the UCLs that would result from assuming normal and lognormal distribution of the data. These

**Highlight 7****EXAMPLE OF DATA TRANSFORMATION AND CALCULATION OF UCL**

This example shows the calculation of a 95 percent UCL of the arithmetic mean concentration for chromium in soil at a Superfund site. This example is applicable only to a scenario in which a spatially random exposure pattern is assumed. The concentrations of chromium obtained from random sampling in soil at this (in mg/kg) are 10, 13, 20, 36, 41, 59, 67, 110, 110, 136, 140, 160, 200, 230, and 1300. Using these data, the following steps are taken to calculate a concentration term for the intake equation:

- (1) Plot the data and inspect the graph. (You may need the help of a statistician for this part, as well as other parts, of the calculation of the UCL.) The plot (not shown, but similar to **Highlight 3**) shows a skew to the right, consistent with a lognormal distribution.
- (2) Transform the data by taking the natural log of the values (i.e., determine  $\ln(x)$ ). For this data set, the transformed values are: 2.30, 2.56, 3.00, 3.58, 3.71, 4.08, 4.20, 4.70, 4.70, 4.91, 4.94, 5.08, 5.30, 5.44, and 7.17.
- (3) Apply the UCL equation in **Highlight 5**, where:

$$\begin{aligned}\bar{x} &= 4.38 \\ s &= 1.25 \\ H &= 3.163 \text{ (based on 95 percent)} \\ n &= 15\end{aligned}$$

The resulting 95 percent UCL of the arithmetic mean is thus found to equal  $e^{(6.218)}$ , or 502 mg/kg.

**Highlight 8****COMPARING UCLs OF THE ARITHMETIC MEAN ASSUMING DIFFERENT DISTRIBUTIONS**

In this example, the data presented in **Highlight 7** are used to demonstrate the difference in the UCL that is seen if the normal distribution approach were inappropriately applied to this data set (i.e., if, in this example, a normal distribution is assumed).

ASSUMED DISTRIBUTION:	Normal	Lognormal
TEST STATISTIC:	Student-t	H- statistic
95 PERCENT UCL (mg/kg):	325	502



examples demonstrate the importance of using the correct assumptions.

### WHERE CAN I GET MORE HELP?

Additional information on Superfund's policy and approach to calculating the concentration term and estimating exposures at waste sites can be obtained in:

- U.S. EPA, *Risk Assessment Guidance for Superfund: Volume I—Human Health Evaluation Manual (Part A)*, EPA/540/1-89/002, December 1989.
- U.S. EPA, *Guidance for Data Usability in Risk Assessment*, EPA/540/G-90/008, (OSWER Directive 9285.7-05), October 1990.
- U.S. EPA, *Risk Assessment Guidance for Superfund (Part A—Baseline Risk Assessment) Supplemental Guidance/Standard Exposure Factors*, OSWER Directive 9285.6-03, May 1991.

Useful statistical guidance can be found in many standard textbooks, including:

- Gilbert, R.O., *Statistical Methods for Environmental Pollution Monitoring*, Van Nostrand Reinhold, New York, New York, 1987.

Questions or comments concerning the concentration term can be directed to:

- Toxics Integration Branch  
Office of Emergency and Remedial Response  
401 M Street, SW.  
Washington, DC 20460  
Phone: 202-260-9486

EPA staff can obtain additional copies of this bulletin by calling EPA's Superfund Document Center at 202-260-9760. Others can obtain copies by contacting NTIS at 703-487-4650.

**K.2 U.S. EPA. (2002) *Calculating the Upper Confidence Limits for Exposure Point Concentrations at Hazardous Waste Sites***

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December 2002

**CALCULATING UPPER CONFIDENCE  
LIMITS FOR EXPOSURE POINT  
CONCENTRATIONS AT HAZARDOUS  
WASTE SITES**

**Office of Emergency and Remedial Response  
U.S. Environmental Protection Agency  
Washington, D.C. 20460**

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**Disclaimer**

This document provides guidance to EPA Regions concerning how the Agency intends to exercise its discretion in implementing one aspect of the CERCLA remedy selection process. The guidance is designed to implement national policy on these issues.

The statutory provisions and EPA regulations described in this document contain legally binding requirements. However, this document does not substitute for those provisions or regulations, nor is it a regulation itself. Thus, it cannot impose legally-binding requirements on EPA, States, or the regulated community, and may not apply to a particular situation based upon the circumstances. Any decisions regarding a particular remedy selection decision will be made based on the statute and regulations, and EPA decisionmakers retain the discretion to adopt approaches on a case-by-case basis that differ from this guidance where appropriate. EPA may change this guidance in the future.

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## 1.0 INTRODUCTION

This document updates a 1992 guidance originally developed to supplement EPA's *Risk Assessment Guidance for Superfund (RAGS), Volume 1 – Human Health Evaluation Manual* (RAGS/HHEM, EPA 1989), which describes a general approach for estimating exposure of individuals to chemicals of potential concern at hazardous waste sites. It addresses a key element of the risk assessment process for hazardous waste sites: estimation of the concentration of a chemical in the environment. This concentration, commonly termed the exposure point concentration (EPC), is a conservative estimate of the average chemical concentration in an environmental medium. The EPC is determined for each individual exposure unit within a site. An exposure unit is the area throughout which a receptor moves and encounters an environmental medium for the duration of the exposure. Unless there is site-specific evidence to the contrary, an individual receptor is assumed to be equally exposed to media within all portions of the exposure unit over the time frame of the risk assessment.

EPA recommends using the average concentration to represent "a reasonable estimate of the concentration likely to be contacted over time" (EPA 1989). The guidance previously issued by EPA in 1992, *Supplemental Guidance to RAGS: Calculating the Concentration Term* (EPA 1992), states that, "because of the uncertainty associated with estimating the true average concentration at a site, the 95 percent upper confidence limit (UCL) of the arithmetic mean should be used for this variable." The 1992 guidance addresses two kinds of data distributions: normal and lognormal. For normal data, EPA recommends an upper confidence limit (UCL) on the mean based on the Student's *t*-statistic. For lognormal data, EPA recommends the Land method using the *H*-statistic. EPA describes approaches for testing distribution assumptions in *Guidance for Data Quality Assessment: Practical Methods for Data Analysis* (EPA 2000b, section 4.2).

The 1992 guidance has been helpful for EPC calculation, but it does not address data distributions that are neither normal nor lognormal. Moreover, as has been widely acknowledged, the Land method can sometimes produce extremely high values for the UCL when the data exhibit high variance and the sample size is small (Singh et al. 1997; Schulz and Griffin 1999). EPA's 1992 guidance recognizes the problem of extremely high UCLs, and recommends that the maximum detected concentration become the default when the calculated UCL exceeds this value. Singh et al. (1997) and Schulz and Griffin (1999) suggest several alternate methods for calculating a UCL for non-normal data distributions. This guidance provides additional tools that risk assessors can use for UCL calculation, and assists in applying these methods at hazardous waste sites. It begins with a discussion of issues related to evaluating the available site data and then presents brief discussions of alternative methods for UCL calculation, with recommendations for their use at hazardous waste sites. In addition, EPA has worked with its contractor, Lockheed Martin to develop a software package, ProUCL, to perform many of the calculations described in this guidance (EPA 2001a). Both ProUCL and this guidance make recommendations for calculating UCLs, and are intended as tools to support risk assessment.

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To obtain a copy of the ProUCL software or receive technical assistance in using it, risk assessors should contact:

Director of the Technical Support Center  
 USEPA Office of Research and Development  
 National Exposure Research Laboratory  
 Environmental Sciences Division  
 Las Vegas, Nevada  
 702-798-2270.

The ultimate responsibility for deciding how best to represent the concentration data for a site lies with the project team.<sup>1</sup> Simply choosing a statistical method that yields a lower UCL is not always the best representation of the concentration data at a site. The project team may elect to use a method that yields a higher (i.e., more conservative) UCL based on its understanding of site-specific conditions, including the representativeness of the data collection process, and the limits of the available statistical methods for calculating a UCL.

## 2.0 APPLICABILITY OF THIS GUIDANCE

This document updates 1992 guidance developed by EPA's Office of Emergency and Remedial Response; yet it can be applied to any hazardous waste site. It provides alternative methods for calculating the 95 percent upper confidence limit of the mean concentration, which can be used at sites subject to the discretion of the regulatory agencies and programs involved. The approaches described in this document are not specific to a particular medium (e.g., soil, groundwater), or receptor (e.g., human ecological), but apply to any media or receptor for which the UCL would be calculated.<sup>2</sup>

This document does not substitute for any statutory provisions or regulations, nor is it a regulation itself. Thus, it cannot impose legally-binding requirements on EPA, States, or the regulatory community, and may not apply to a particular situation based upon the circumstances. Any decision regarding cleanup of a particular site will be made based on the statutes and regulations, and EPA decisionmakers retain the discretion to adopt approaches on a case-by-case basis that differ from this guidance to a particular situation. The Agency accepts public input on this document at any time.

This guidance is based on the state of knowledge at present. The practices discussed herein may be refined, updated, or superseded by future advances in science and mathematics.

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<sup>1</sup> The project team typically consists of a site manager (e.g., the Remedial Project Manager) and a multidisciplinary team of technical experts, including human health and ecological risk assessors, hydrogeologists, chemists, toxicologists, and quality assurance specialists.

<sup>2</sup> Note that this guidance does not apply to lead-contaminated sites. The Technical Review Working Group for Lead recommends that the average concentration is used in evaluating lead exposures (see <http://www.epa.gov/superfund/programs/lead/trwhome.htm>).

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### 3.0 DATA EVALUATION

In the risk assessment process, data evaluation precedes exposure assessment. Because this guidance deals with a component of exposure assessment, it therefore assumes that data have already undergone validation and evaluation and that the data have been determined to meet data quality objectives (DQOs) for the project in question. DQOs are important for any project where environmental data are used to support decision-making, as at hazardous waste sites.

One factor to consider in data evaluation is whether the number of sample measurements is sufficient to characterize the site or exposure unit. The minimum number of samples to conduct any of the statistical tests described in this document should be determined using the DQO process (EPA 2000a). Use of the methods described in this guidance is not a substitute for obtaining an adequate number of samples. Sample size is especially important when there is large variability in the underlying distribution of concentrations. However, defaulting to the maximum value of small data sets may still be the last resort when the UCL appears to exceed the range of concentrations detected.

Another important issue to consider is the method of sampling. All the statistical methods described in this guidance for calculating UCLs are based on the assumption of random sampling. At many hazardous waste sites, however, sampling is focused on areas of suspected contamination. In such cases, it is important to avoid introducing bias into statistical analyses. This can be achieved through stratified random sampling, i.e., random sampling within specified targeted areas. So long as the statistical analysis is constructed properly (i.e., there is no mixing of samples across different populations) bias can be minimized. The risk assessor should always note any potential bias in EPC estimates.

The risk assessor should also consider the duration of exposure and the time scale of the toxicity. For example, a chronic exposure may warrant the use of different concentrations or sample locations from an acute exposure. The time periods over which data were collected should also be considered. See EPA 1989, Chapters 5.1 and 6.4.2, for further details.

Once a set of data from a site has been evaluated and validated, it is appropriate to conduct exploratory analysis to determine whether there are outliers or a substantial number of non-detect values that can adversely affect the outcome of statistical analyses. The following sections describe the potential impact of outliers and non-detect values on the calculation of UCLs and approaches for addressing these types of values.

#### **3.1 Outliers**

Outliers are values in a data set that are not representative of the set as a whole, usually because they are very large relative to the rest of the data. There are a variety of statistical tests for determining whether one or more observations are outliers (EPA 2000b, section 4.4). These tests should be used judiciously, however. It is common that the distribution of concentration data at a site is strongly skewed so that it contains a few very high values corresponding to local hot spots of contamination. The receptor could be exposed to these hot spots, and to estimate the EPC correctly it is important to take account of these values. Therefore, one should be careful not to exclude values merely because they are large relative to the rest of the data set.



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Extreme values in the data set may represent true spatial variation in concentrations. If an observation or group of observations is suspected to be part of a different contamination source or exposure unit, then regrouping of the data may be most appropriate. In this case, it may be necessary to evaluate these data as a separate hot spot or to resample. The behavior of the receptor and the size and location of the exposure unit will determine which sample locations to include. Such decisions depend on project-specific assessments based on the conceptual site model.

EPA guidance suggests that, when outliers are suspected of being unreliable and statistical tests show them to be unrepresentative of the underlying data set, any subsequent statistical analyses should be conducted both with and without the outlier(s) (EPA 2000b). In addition, the entire process, including identification, statistical testing and review of outliers, should be fully documented in the risk characterization.

### **3.2 Non-detects**

Chemical analyses of contaminant concentrations often result in some samples being reported as below the sample detection limit (DL). Such values are called non-detects. Non-detects may correspond to concentrations that are actually or virtually zero, or they may correspond to values that are considerably larger than zero but which are below the laboratory's ability to provide a reliable measurement. Elevated detection limits need to be investigated, especially if there are high percentages of non-detects. It is not appropriate to simply account for elevated detection limits with statistical techniques; improvements in sampling and analysis methods may be needed to lower detection limits.

In this guidance, the term "detection limit" is used to represent the reported limit of the non-detect. In reality, this could be any of a number of detection or quantitation limits. For further discussion of detection and quantitation limits in the risk assessment, see text box and Chapter 5 of EPA 1989.

#### **Alternative Quantitation Limits**

**Method Detection Limit (MDL):** The lowest concentration of a hazardous substance that a method can detect reliably in either a sample or blank.

**Contract-Required Quantitation Limit (CRQL):** The substance-specific level that a CLP laboratory must be able to routinely and reliably detect in specific sample matrices. The CRQL is not the lowest detectable level achievable, but rather the level that a CLP laboratory must reliably quantify. The CRQL may or may not be equal to the quantitation limit of a given substance in a given sample.

Source: Superfund Glossary of Terms and Acronyms (<http://www.epa.gov/superfund/resources/hrstrain/htmain/glossal.htm>)

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In the statistical literature, data sets containing non-detects are called censored or left-censored. The detection limit achieved for a particular sample depends on the sensitivity of the measuring method used, the instrument quantitation limit, and the nature of dilutions and other preparations employed for the sample. In addition, there may be different degrees of censoring. For instance, some laboratories use the letter code "J" to indicate that a value was below the quantitation limit and the letter "U" to indicate that a value was below the detection limit. These code systems vary among laboratories, however, and it is essential to understand what the laboratory notations indicate about the reliability of its measurements.<sup>3</sup> Censoring can cause problems in calculating the UCL. There are several common options for handling non-detects.

**Reexamining the conceptual site model** may suggest that the data be partitioned. For instance, it may be clear from the spatial pattern of non-detects in the data that the region sampled can be subdivided into contaminated and non-contaminated areas. Evidence for this depends on the observed pattern of contamination, how the contamination came to be located in the medium, and how the receptors will come in contact with the medium. It may be necessary to collect more samples to obtain an adequate site characterization.

**Simple Substitution methods** assign a constant value or constant fraction of the detection limit (DL) to the non-detects. Three common conventions are: (1) assume non-detects are equal to zero; (2) assume non-detects are equal to the DL; or (3) assume non-detects are equal to one-half the DL. Whatever proxy value is assigned, it is then used as though it were the reliably estimated value for that measurement. Because of the complicated formulas used to compute UCLs, there is no general rule about which substitution rule will yield an appropriate UCL. The uncertainty associated with the substitution method increases, and its appropriateness decreases, as the detection limit becomes larger and as the number of non-detects in the data set increases.

**Bounding methods** estimate limits on the UCL in a distribution-free way. This method involves determining the lower and upper bounds of the UCL based on the full range of possible values for non-detects. If the uncertainty arising from censoring is relatively small, then the difference between the lower and upper bound estimates will be small. It is not possible to bound the UCL by using simple substitution methods such as computing the UCL once with the non-detects replaced by zeros and once with the non-detects replaced by their respective detection limits. Sometimes using all zeros will inflate the estimate of the standard deviation of the concentration values to such a degree that the resulting value for the UCL is larger than the value from using the detection limits (Ferson et al. 2002, Rowe 1988, Smith 1995). See Appendix A for an example of how to compute bounds on the UCL.

**Distributional methods** rely on applying an assumption that the shape of the distribution of non-detect values is similar to that of measured concentrations above the detection limit. EPA provides guidance on handling non-detects using several distributional methods, including Cohen's method (EPA 2000b, section 4.7). In addition, Helsel (1990) reviews a variety of distributional methods (see also Hass and Scheff 1990; Gleit 1985; Kushner 1976; Singh and Nocerino 2001). *EnvironmentalStats for S-PLUS* (Millard 1997) offers an array of methods for estimating parameters from censored data sets.

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<sup>3</sup> Information concerning the quantitation limits also should be incorporated into the appropriate supplemental tables in the framework for risk assessment planning, reporting, and review described in the *Risk Assessment Guidance for Superfund Volume 1: Human Health Evaluation Part D (RAGS, Part D)* (EPA 1998.)

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The appropriate method to use depends on the severity of the censoring, the size of the data set, and what distributional assumptions are reasonable. There are five recommendations about how to treat censoring in the estimation of UCLs.

- 1) Detection limits should always be reported for non-detects. Non-detects should also be reported with observed values where possible.
- 2) It is inappropriate to convert non-detects into zeros without specific justification (e.g., the analyte was not detected above the detection limit in any sample at the site).
- 3) If a bounding analysis reveals that the quantitative effects of censoring are negligible, then no further analysis may be required.
- 4) If further analysis is desired, consider using a distribution-specific method.
- 5) If the proportion of non-detects is high ( $\geq 75\%$ ) or the number of samples is small ( $n < 5$ ), no method will work well. In this case, it is reasonable to report the percentage of data below the detection limit, and resort again to a bounding approach in which non-detects are replaced by the detection limit and used to compute a UCL value that is reported as a number likely to be considerably larger than the true mean.

#### 4.0 UCL CALCULATION METHODS

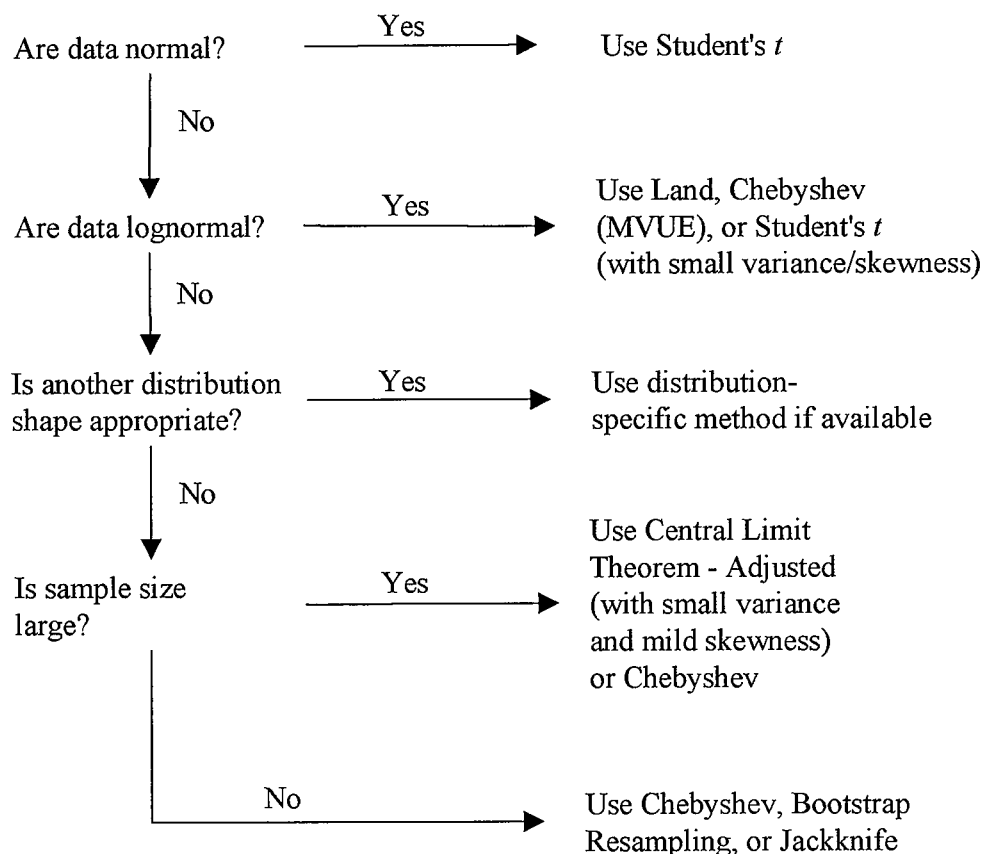
There are a number of different methods for calculating UCLs. Before an appropriate method can be selected the site data must be characterized through exploratory analysis. Fitting distributions to the data is a crucial part of this exploratory data analysis (Schulz and Griffin 1999). As recommended by EPA (1992), "where there is a question about the distribution of the data set, a statistical test should be used to identify the best distributional assumption for the data set." This is necessary because no single distribution type fits all environmental data sets. Risk assessors deal with some environmental data sets that appear normally distributed, and with others that appear lognormally distributed. They also encounter data sets that do not fit either normal or lognormal distributions. Distributions can be analyzed by a variety of methods, many of which are described in Gilbert (1987) and EPA (2000b). Data plotting can also help identify a useful distributional assumption. Some of these methods have been incorporated in the ProUCL software. Whatever method is used, it should be chosen in consultation with the EPA regional risk assessor and other project team members as appropriate. The assistance of a statistician may also be helpful in some cases.

The two most commonly used methods for computing UCLs are distributional methods. When the concentration distribution is normal, the classical approach based on the Student's *t*-statistic has typically been used. When the distribution is lognormal, the Land method based on the *H*-statistic has been used. Distribution-free or nonparametric methods are available if the risk assessor cannot reasonably make assumptions about the distributional type. EPA describes several methods (EPA, 2000c). For large data sets, an approach based on the Central Limit Theorem with a correction for positive skewness may be used. For data sets that are not large enough for this approach, there is more than one approach available, although none is ideal in all circumstances. General methods include an approach based on the Chebyshev inequality and an approach based on the bootstrap resampling procedure. These are described in EPA (2000c) and in Schulz and Griffin (1999). Both papers give examples and comparisons of the UCLs calculated by various methods. The flow chart shown in Figure 1 summarizes the recommendations in this guidance.

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It should be noted that the “variance” in Figure 1 represents the variance of the log-transformed data. For detailed definitions of skewness, refer to the User’s Guide for the ProUCL software.

**Figure 1: UCL Method Flow Chart**



Risk assessors are encouraged to use the most appropriate estimate for the EPC given the available data. The flow chart in Figure 1 provides general guidelines for selecting a UCL calculation method. This guidance presents descriptions of these methods, including their applicability, advantages and disadvantages. It also includes examples of how to calculate UCLs using the methods. While the methods identified in this guidance may be useful in many situations, they will probably not be appropriate for all hazardous waste sites. Moreover, other methods not specifically described in this guidance may be most appropriate for particular sites. The EPA risk assessor should be involved in the decision of which method(s) to use.

#### **4.1 UCL Calculation with Methods for Specific Distributions**

This section of the guidance presents methods for calculating UCLs when data can be shown to fit a specific distribution. Directions for using methods to calculate UCL for normal, lognormal, and other specific distributions are included, as are example calculations.

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**UCLs for Normal Distributions**

If the data are normally distributed, then the one-sided  $(1-\alpha)$  upper confidence limit  $UCL_{1-\alpha}$  on the mean should be computed in the classical way using the Student's  $t$ -statistic (EPA 1992; Gilbert 1987, page 139; Student 1908). There is no change in EPA's prior recommendations for this type of data set (EPA 1992). Exhibit 1 gives the procedure for computing the UCL of the mean when the underlying distribution is normal. Exhibit 2 gives a numerical example of an application of the method.

**Exhibit 1: Directions for Computing UCL for the Mean of a Normal Distribution —  
Student's  $t$**

Let  $X_1, X_2, \dots, X_n$  represent the  $n$  randomly sampled concentrations.

STEP 1: Compute the sample mean  $\bar{X} = \frac{1}{n} \sum_{i=1}^n X_i$ .

STEP 2: Compute the sample standard deviation  $s = \sqrt{\frac{1}{n-1} \sum_{i=1}^n (X_i - \bar{X})^2}$ .

STEP 3: Use a table of quantiles of the Student's  $t$  distribution to find the  $(1-\alpha)^{\text{th}}$  quantile of the Student's  $t$  distribution with  $n-1$  degrees of freedom. For example, the value at the 0.05 level with 40 degrees of freedom is 1.684. A table of Student's  $t$  values can be found in Gilbert (1987, page 255, where the values are indexed by  $p=1-\alpha$ , rather than  $\alpha$  level). The  $t$  value appropriate for computing the 95% UCL can be obtained in Microsoft Excel® with the formula  $\text{TINV}((1-0.95)*2, n-1)$ .

STEP 4: Compute the one-sided  $(1-\alpha)$  upper confidence limit on the mean

$$UCL_{1-\alpha} = \bar{X} + t_{\alpha, n-1} s / \sqrt{n}$$

**Exhibit 2: An Example Computation of UCL for a Normal Distribution — Student's  $t$** 

25 samples were collected at random from an exposure unit. The values observed are 228, 552, 645, 208, 755, 553, 674, 151, 251, 315, 731, 466, 261, 240, 411, 368, 492, 302, 438, 751, 304, 368, 376, 634, and 810  $\mu\text{g/L}$ . It seems reasonable that the data are normally distributed, and the Shapiro-Wilk  $W$  test for normality fails to reject the hypothesis that they are ( $W = 0.937$ ). The UCL based on Student's  $t$  is computed as follows.

STEP 1: The sample mean of the  $n=25$  values is  $\bar{X} = 451$ .

STEP 2: The sample standard deviation of the values is  $s = 198$ .

STEP 3: The  $t$ -value at the 0.05 level for 25-1 degrees of freedom is  $t_{0.05,25-1} = 1.710$ .

STEP 4: The one-sided 95% upper confidence limit on the mean is therefore

$$UCL_{95\%} = 451 + 1.710 \times 198 / \sqrt{25} = 519$$

*Testing for normality.* For mildly skewed data sets, the student's  $t$ -statistic approach may be used to compute the UCL of the mean. But for moderate to highly skewed data sets, the  $t$ -statistic-based UCL can fail to provide the specific coverage for the population mean. This is especially true for small  $n$ . For instance, the 95% UCL based on 10 random samples from a lognormal distribution with mean 4.48 and standard deviation 5.87 will underestimate the true mean about 20% of the time, rather than the nominal rate of 5%. Therefore it is important to test the data for normality. EPA (2000b, section 4.2) gives guidance for several approaches for testing normality. The tests described therein are available in DataQUEST and ProUCL, which are convenient software tools (EPA 1997 and 2001a).

*Accounting for non-detects.* The use of substitution methods to account for non-detects is recommended only when a very small percentage of the data is censored (e.g.,  $\bullet$  15%), under the presumption that the numerical consequences of censoring will be minor in this case. As the percentage of the data censored increases, substitution methods tend to alter the distribution and violate the assumption of normality. Moreover, the effect of the various substitution rules on UCL estimation is difficult to predict. Replacing non-detects with half the detection limit can underestimate the UCL, and replacing them with zeros may overestimate the UCL (because doing so inflates the estimate of the standard deviation).

When censoring is moderate (e.g.,  $>15\%$  and  $\bullet$  50%), it is preferable to account for non-detects with Cohen's method (Gilbert 1987). EPA provides guidance on the use of Cohen's method, which is a maximum likelihood method for correcting the estimates of the sample mean and the sample variance to account for the presence of non-detects among the data (EPA 2000b, beginning on page 4-43). This method requires that the detection limit be the same for all the data and that the underlying data are normally distributed.

**UCLs for Lognormal Distributions**

It is inappropriate to extend the methods of the previous section to lognormally distributed samples by log-transforming the data, computing a UCL and then back-transforming the results. For

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concentration data sets that appear to be lognormally distributed, it may instead be preferable to use one of several methods available that are specifically well-suited to this type of distribution. These methods are described in the following sections.

### **Land Method**

In past guidance, EPA had recommended using the Land method to compute the upper confidence limit on the mean for lognormally distributed data (Land 1971, 1975; Gilbert 1987; EPA 1992; Singh et al. 1997). This method requires the use of the  $H$ -statistic, tables for which were published by Land (1975) and Gilbert (1987, Tables A10 and A12). Exhibit 3 gives step-by-step directions for this method and Exhibit 4 gives a numerical example of its application.

*Caveats about this method.* Land's approach is known to be sensitive to deviations from lognormality. The formula may commonly yield estimated UCLs substantially larger than necessary when distributions are not truly lognormal if variance or skewness is large (Gilbert 1987). When sample sizes are small (less than 30), the method can be impractical even when the underlying distribution is lognormal (Singh et al. 1997).

#### **Exhibit 3: Directions for Computing UCL for the Mean of a Lognormal Distribution— Land Method**

Let  $X_1, X_2, \dots, X_n$  represent the  $n$  randomly sampled concentrations.

STEP 1: Compute the arithmetic mean of the log-transformed data  $\overline{\ln X} = \frac{1}{n} \sum_{i=1}^n \ln(X_i)$ .

STEP 2: Compute the associated standard deviation  $s_{\ln X} = \sqrt{\frac{1}{n-1} \sum_{i=1}^n (\ln(X_i) - \overline{\ln X})^2}$ .

STEP 3: Look up the  $H_{1-\alpha}$  statistic for sample size  $n$  and the observed standard deviation of the log-transformed data. Tables of these values are given by Gilbert (1987, Tables A-10 and A-12) and Land (1975).

STEP 4: Compute the one-sided  $(1-\alpha)$  upper confidence limit on the mean

$$UCL_{1-\alpha} = \exp \left( \overline{\ln X} + s_{\ln X}^2 / 2 + H_{1-\alpha} s_{\ln X} / \sqrt{n-1} \right)$$

*Testing for lognormality.* Because the Land method assumes lognormality, it is very important to test this assumption. EPA gives guidance for several approaches to testing distribution assumptions (EPA 2000b, section 4.2). The tests are also available in the DataQUEST and ProUCL software tools (EPA 1997 and 2001a).

**Exhibit 4: An Example Computation of UCL for a Lognormal Distribution —  
Land Method**

31 samples were collected at random from an exposure unit. The observed values are 2.8, 22.9, 3.3, 4.6, 8.7, 30.4, 12.2, 2.5, 5.7, 26.3, 5.4, 6.1, 5.2, 1.8, 7.2, 3.4, 12.4, 0.8, 10.3, 11.4, 38.2, 5.6, 14.1, 12.3, 6.8, 3.3, 5.2, 2.1, 19.7, 3.9, and 2.8 mg/kg. Because of their skewness, the data may be lognormally distributed. The Shapiro-Wilk  $W$  test for normality rejects the hypothesis, at both the 0.05 and 0.01 levels, that the distribution is normal. The same test fails to reject at either level the hypothesis that the distribution is lognormal. The UCL on the mean based on Land's  $H$  statistic is computed as follows.

STEP 1: Compute the arithmetic average of the log-transformed data  $\overline{\ln X} = 1.8797$ .

STEP 2. Compute the standard deviation of the log-transformed data  $s_{\ln X} = 0.8995$ .

STEP 3. The  $H$  statistic for  $n = 31$  and  $s_{\ln X} = 0.90$  is 2.31.

STEP 4: The one-sided 95% upper confidence limit on the mean is therefore

$$UCL_{95\%} = \exp\left(1.8797 + 0.8995^2 / 2 + 2.31 \times 0.8995 / \sqrt{31-1}\right) = 14.4$$

*Accounting for non-detects.* Gilbert (1987, page 182) suggests extending Cohen's method to account for non-detect values in lognormally distributed concentrations. Cohen's method (EPA 2000b, page 4-43) assumes the data are normally distributed, so it must be applied to the log-transformed concentration values. If  $\hat{\mu}_y$  and  $\hat{\sigma}_y$  are the corrected sample mean and standard deviation, respectively, of the log-transformed concentrations, then the corrected estimates of the mean and standard deviation of the underlying lognormal distribution can be obtained from the following expressions:

$$\hat{\mu} = \exp(\hat{\mu}_y + \hat{\sigma}_y^2 / 2)$$

$$\hat{\sigma} = \hat{\mu} \sqrt{\exp(\hat{\sigma}_y^2) - 1}$$

This method requires there be a single detection level for all the data values.

**Chebyshev Inequality Method**

Singh et al. (1997) and EPA (2000c) suggest the use of the Chebyshev inequality to estimate UCLs which should be appropriate for a variety of distributions so long as the skewness is not very large. The one-sided version of the Chebyshev inequality (Allen 1990, page 79; Savage 1961, page 216) is appropriate in this context (cf. Singh et al. 1997, EPA 2000c). It can be applied to the sample mean to obtain a distribution-free estimate of the UCL for the population mean when the population variance or standard deviation are known. In practice, however, these values are not known and must be estimated from data. For lognormally distributed data sets, Singh et al. (1997) and EPA (2000c) suggest using the minimum-variance unbiased estimators (MVUE) for the mean and variance to obtain an UCL of the mean. (See also Gilbert 1987, for discussion of the MVUE). This



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approach may yield an estimated UCL that is more useful than that obtained from the Land method (when the underlying distribution of concentrations is lognormal). This alternative approach for a lognormal distribution is described in Exhibit 5 and is available in the ProUCL software tool (EPA 2001a). A numerical illustration of the Chebyshev inequality method using the sample mean and standard deviation appears in Exhibit 6. In this example the estimate of the UCL based on the Chebyshev inequality is less than that based on the Land method. The Chebyshev inequality estimate of the UCL is 1,965 mg/kg; while applying the Land method to this same data set yields a higher UCL estimate of 2,658 mg/kg.

**Exhibit 5: Steps for UCL Calculation Based on the Chebyshev Inequality — MVUE Approach for Lognormal Distributions**

Let  $X_1, X_2, \dots, X_n$  represent the  $n$  randomly sampled concentrations.

STEP 1: Compute the arithmetic mean of the log-transformed data  $\overline{\ln X} = \frac{1}{n} \sum_{i=1}^n \ln(X_i)$ .

STEP 2: Compute the associated variance  $s_{\ln X}^2 = \frac{1}{n-1} \sum_{i=1}^n (\ln(X_i) - \bar{y})^2$ .

STEP 3: Compute the minimum-variance unbiased estimator (MVUE) of the population mean for a lognormal distribution  $\hat{\mu}_{LN} = \exp(\overline{\ln X}) g_n(s_{\ln X}^2/2)$ , where  $g_n$  denotes a function for which tables are available (Aitchison and Brown 1969, Table A2; Koch and Link 1980, Table A7).

STEP 4: Compute the MVUE of the associated variance of this mean

$$\sigma_{\mu}^2 = \exp(2\overline{\ln X}) \left( \left( g_n(s_{\ln X}^2/2) \right)^2 - g_n\left(\frac{n-2}{n-1} s_{\ln X}^2\right) \right)$$

STEP 5: Compute the one-sided  $(1-\alpha)$  upper confidence limit on the mean

$$UCL_{1-\alpha} = \hat{\mu}_{LN} + \sqrt{\left(\frac{1}{\alpha} - 1\right) \sigma_{\mu}^2}$$

*Caveats about the Chebyshev method.* EPA (2000c) points out that for highly skewed lognormal data with small sample size and large standard deviation, the Chebyshev 99% UCL may be more appropriate than the 95% UCL, because the Chebyshev 95% UCL may not provide adequate coverage of the mean. As skewness increases further, the Chebyshev method is not recommended. See the ProUCL User's Guide (2001a) for specific recommendations on use of these two UCL estimates.

**Exhibit 6: An Example Computation of UCL Based on the Chebyshev Inequality**

29 samples were collected at random from an exposure unit. The observed values are 107, 175, 1796, 2002, 109, 30, 273, 83, 127, 254, 466, 12, 403, 31, 1042, 923, 24, 537, 5667, 59, 158, 59, 353, 10, 8, 33, 1129, 3 and 279 mg/kg. The observed skewness of this data set is 3.8, and these data may be lognormally distributed. The assumption of normality is rejected at the 0.05 level by a Shapiro-Wilk W test, but the same test fails to reject a test of lognormality even at the 0.1 level. The UCL on the mean can be computed based on the Chebyshev Inequality as follows.

- STEP 1: The arithmetic mean of the log-transformed data  $\overline{\ln X}$  is 4.9690.
- STEP 2: The associated variance  $S_{\ln X}^2 = 3.3389$ .
- STEP 3: The MVUE of the mean for a lognormal distribution  $\hat{\mu}_{LN} = 666.95$ .
- STEP 4: The MVUE of the variance of the mean  $\sigma_{\mu}^2 = 88552$ .
- STEP 5: The resulting one-sided 95% upper confidence limit on the mean of the concentration

$$UCL_{95\%} = 666.95 + \sqrt{(19)88552} = 1,965$$

The 95% UCL based on the Land method for these data would be 2,658.

EPA (2000c, Table 7) suggests that the Chebyshev inequality method for computing the UCL may be preferred over the Land method, even for lognormal distributions, in certain situations. Exhibit 7 describes the conditions, in terms of the sample size and the standard deviation of the log-transformed data, under which the Chebyshev inequality method will probably yield more useful results than the Land method.

<b>Exhibit 7</b> <b>Conditions Likely to Favor Use of Chebyshev Inequality (MVUE)</b> <b>over Land Method</b>		
<b>Standard deviation of log-transformed data</b>	<b>Sample Size</b>	<b>Recommendation</b>
1 - 1.5	<25	95% Chebyshev (MVUE) UCL
1.5 - 2	<20	99% Chebyshev (MVUE) UCL
	20 - <50	95% Chebyshev (MVUE) UCL
2 - 2.5	<25	99% Chebyshev (MVUE) UCL
	25 - 70	95% Chebyshev (MVUE) UCL
2.5 - 3.0	<30	99% Chebyshev (MVUE) UCL
	30 - <70	95% Chebyshev (MVUE) UCL

#### UCLs for Other Specific Distribution Types

Methods for computing UCLs on the mean of other types of distributions have appeared in the statistical literature. For example, Johnson (1978) describe a method for computing the UCL for asymmetrical distributions such as the exponential. Schulz and Griffin (1999) described Wong's (1993) method for obtaining confidence limits on the mean of a gamma distribution. In general, if there are arguments that suggest a population of concentrations should fit a particular distribution shape, and if statistical testing confirms the expected shape reasonably conforms with available data, then the UCL computed by a method developed specifically for the distribution shape, if one exists, is likely to be appropriate for the data set. An analyst should consider using a distribution-specific method if possible because it is likely to produce more valid statistical results. The advice and support of a statistician may be invaluable in such cases, both for characterizing the distribution and for identifying and evaluating possible ways to derive confidence limits.

#### 4.2 UCL Calculation With Nonparametric or Distribution-free Methods

There are also distribution-free approaches to computing UCLs on the mean that do not make specific assumptions about the shape of the underlying distribution of concentrations. While these methods assume the samples are representative of the underlying distribution of concentrations, they require no assumptions about the shape of that distribution and are applicable to a variety of situations. Although parametric statistical methods that depend on a distributional assumption are usually more efficient and powerful than nonparametric methods, it can be difficult to justify their use through empirical testing of the shape of the distribution. In such cases, one of the following nonparametric, or distribution-free techniques are often preferred. For information on how to account for non-detects, see the earlier discussion under "Data Evaluation" above.

**Central Limit Theorem (Adjusted)**

If sample size is sufficiently large, the Central Limit Theorem (CLT) implies that the mean will be normally distributed, no matter how complex the underlying distribution of concentrations might be. This is the case even if the underlying distribution is strongly skewed, has outliers, or is a mixture of different populations, so long as it is stationary (not changing over time), has finite variance, and the samples are collected independently and randomly. However, the theorem does not say how many samples are sufficient for normality to hold. When sample size is moderate or small the means will not generally be normally distributed, and this non-normality is intensified by the skewness of the underlying distribution. Chen (1995) suggested an approach that accounts for positive skewness. Singh et al. (1997) and EPA (2000c) call this approach the “adjusted CLT” method. They suggest it is an appropriate alternative to the distribution-specific Land’s method even if the distribution is lognormal when the standard deviation is less than one and sample size is larger than 100. Exhibit 8 describes the steps for this method, and Exhibit 9 gives a numerical example.

**Exhibit 8: Directions for Computing UCL Using the Central Limit Theorem (Adjusted)**

Let  $X_1, X_2, \dots, X_n$  represent the  $n$  randomly sampled concentrations.

STEP 1: Compute the sample mean  $\bar{X} = \frac{1}{n} \sum_{i=1}^n X_i$ .

STEP 2: Compute the sample standard deviation  $s = \sqrt{\frac{1}{n-1} \sum_{i=1}^n (X_i - \bar{X})^2}$ .

STEP 3: Compute the sample skewness  $\beta = \frac{n}{(n-1)(n-2)} \sum_{i=1}^n \left( \frac{x_i - \bar{x}}{s} \right)^3$ . This can be calculated in Microsoft® Excel with the SKEW function.

STEP 4: Let  $z_\alpha$  be the  $(1-\alpha)^{\text{th}}$  quantile of the standard normal distribution. For the 95% confidence level,  $z_\alpha = 1.645$ .

STEP 5: Compute the one-sided  $(1-\alpha)$  upper confidence limit on the mean

$$UCL_{1-\alpha} = \bar{X} + \left( z_\alpha + \frac{\beta}{6\sqrt{n}} (1 + 2z_\alpha^2) \right) s / \sqrt{n}.$$

**Exhibit 9: Example UCL Computation Based on the Central Limit Theorem (Adjusted)**

60 samples were collected at random from an exposure unit. The values observed are 35, 111, 105, 27, 25, 20, 17, 21, 32, 32, 23, 17, 35, 32, 29, 25, 97, 20, 26, 18, 17, 18, 26, 25, 16, 28, 29, 28, 21, 119, 23, 98, 20, 21, 24, 21, 22, 117, 27, 25, 22, 21, 26, 24, 33, 33, 21, 24, 30, 31, 23, 30, 28, 25, 22, 23, 25, 28, 26, and 107 mg/L. Filliben's test shows that this distribution is significantly different (at the 1% level) from both a normal and a lognormal distribution. The UCL based on the Central Limit Theorem is computed as follows.

STEP 1: The sample mean of the  $n=60$  values is  $\bar{X} = 34.57$ .

STEP 2: The sample standard deviation of the values is  $s = 27.33$ .

STEP 3: The sample skewness  $\beta = 2.366$ .

STEP 4: The  $z$  statistic is 1.645.

STEP 5: The one-sided 95% upper confidence limit on the mean is

$$UCL_{95\%} = 34.57 + \left( 1.645 + \frac{2.366}{6\sqrt{60}} (1 + 2 \times 1.645^2) \right) 27.33 / \sqrt{60} = 42$$

*Caveats about this method.* A sample size of 30 is sometimes prescribed as sufficient for using an approach based on the Central Limit Theorem, but when using this CLT or adjusted CLT method and the data are skewed (as many concentration data sets are), larger samples may be needed to approximate normality. EPA's ProUCL User's Guide (2001) suggests that a sample size of 100 or more may be needed, based on Monte Carlo studies by EPA (2000c).

**Bootstrap Resampling**

Bootstrap procedures (Efron 1982) are robust nonparametric statistical methods that can be used to construct approximate confidence limits for the population mean. In these procedures, repeated samples of size  $n$  are drawn with replacement from a given set of observations. The process is repeated a large number of times (e.g., thousands), and each time an estimate of the desired unknown parameter (e.g., the sample mean) is computed. There are different variations of the bootstrap procedure available. One of these, the bootstrap  $t$  procedure, is described in the ProUCL User's Guide (EPA 2001a). An elaborated bootstrap procedure that takes bias and skewness into account is described in Exhibit 10 (Hall 1988 and 1992; Manly 1997; Schulz and Griffin 1999; Zhou and Gao 2000).

*Caveats about resampling.* Bootstrap procedures assume only that the sample data are representative of the underlying population. However, since they involve extensive resampling of the data and, thus, exploit more of the information in a sample, that sample must be a statistically accurate characterization of the underlying population in all respects (not just in its mean and standard deviation). In practice, it is random sampling that satisfies the representativeness assumption. Therefore the data must be random samples of the underlying population. Bootstrapping procedures are inappropriate for use with data that were idiosyncratically collected or focused especially on contamination hot spots.

**Exhibit 10: Steps for Calculating a Hall's Bootstrap Estimate of UCL**

Let  $X_1, X_2, \dots, X_n$  represent the  $n$  randomly sampled concentrations.

STEP 1: Compute the sample mean  $\bar{X} = \frac{1}{n} \sum_{i=1}^n X_i$ .

STEP 2: Compute the sample standard deviation  $s = \sqrt{\frac{1}{n} \sum_{i=1}^n (X_i - \bar{X})^2}$ .

STEP 3: Compute the sample skewness  $k = \frac{1}{ns^3} \sum_{i=1}^n (X_i - \bar{X})^3$ .

STEP 4: For  $b = 1$  to  $B$  (a very large number) do the following:  
 4.1: Generate a bootstrap sample data set; i.e., for  $i = 1$  to  $n$  let  $j$  be a random integer between 1 and  $n$  and add observation  $X_j$  to the bootstrap sample data set.  
 4.2: Compute the arithmetic mean  $\bar{X}_b$  of the data set constructed in step 4.1.  
 4.3: Compute the associated standard deviation  $s_b$  of the constructed data set.  
 4.4: Compute the skewness  $k_b$  of the constructed data using the formula in Step 3.  
 4.5: Compute the studentized mean  $W = (\bar{X}_b - \bar{X}) / s_b$ .  
 4.6: Compute Hall's statistic  $Q = W + k_b W^2 / 3 + k_b^2 W^3 / 27 + k_b / (6n)$ .

STEP 5: Sort all the  $Q$  values computed in Step 4 and select the lower  $\alpha^{\text{th}}$  quantile of these  $B$  values. It is the  $(\alpha B)^{\text{th}}$  value in an ascending list of  $Q$ 's. This value is from the *left* tail of the distribution.

STEP 6: Compute  $W(Q) = \frac{3}{k} \left( \left( 1 + k \left( Q_\alpha - \frac{k}{6n} \right) \right)^{1/3} - 1 \right)$ .

STEP 7: Compute the one-sided  $(1-\alpha)$  confidence limit on the mean.

$$UCL_{1-\alpha} = \bar{X} - W(Q_\alpha)s$$

**Exhibit 11: An Example Computation of Bootstrap Estimate of UCL**

Using the same concentration values given in Exhibit 4, the UCL can also be computed based on the Bootstrap Resampling method.

STEP 1: The sample mean of the  $n = 31$  values is  $\bar{X} = 9.59$ .

STEP 2: The standard deviation (using  $n$  as divisor) of the values is  $s = 8.946$ .

STEP 3: The skewness  $k = 1.648$ .

The Pascal-language software shown in Appendix B estimates the UCL with 100,000 bootstrap iterations. The one-sided 95% UCL on the mean is 13.3. Because this value depends on random deviates, it can vary slightly on recalculation.

**Jackknife Procedure**

Like bootstrap, the jackknife technique is a robust procedure based on resampling (Tukey 1977). In this procedure repeated samples are drawn from a given set of observations by omitting each observation in turn, yielding  $n$  data sets of size  $n-1$ . An estimate of the desired unknown parameter (e.g., sample mean) is then computed for each sample. When the standard estimators are used for the mean and standard deviation, this procedure reduces to the UCL based on Student's  $t$ . However, when other estimators (such as MVUE) are used this jackknife procedure does not reduce to the UCL based on Student's  $t$ . Singh et al. (1997) suggest that this method could be used with other estimators for the population mean and standard deviation to yield UCLs that may be appropriate for a variety of distributions.

**Chebyshev Inequality Method**

As described previously, Singh et al. (1997) and EPA (2000c) suggested the use of the Chebyshev inequality to estimate UCLs which should be appropriate for a variety of distributions as long as the skewness is not very large. The one-sided version of the Chebyshev inequality (Allen 1990, page 79; Savage 1961, page 216) is appropriate in this context (cf. Singh et al. 1997, EPA 2000c). It can be applied to the sample mean to obtain a distribution-free estimate of the UCL for the population mean when the population variance or standard deviation are known. In practice, however, these values are not known and must be estimated from data. Singh et al. (1997) and EPA (2000c) suggest that the population mean and standard deviation can be estimated by the sample mean and sample standard deviation. This approach is described in Exhibit 12 and is available in the ProUCL software tool (EPA 2001a). A numerical illustration of the Chebyshev inequality method using the sample mean and standard deviation appears in Exhibit 13.

*Caveats about the Chebyshev method.* Although the Chebyshev inequality method makes no distributional assumptions, it does assume that the parametric standard deviation of the underlying distribution is known. As Singh et al. (1997) acknowledge, when this parameter must be estimated from data, the estimate of the UCL is not guaranteed to be larger than the true mean with the prescribed frequency implied by the  $\alpha$  level. In fact, using only an estimate of the standard deviation can substantially underestimate the UCL when the variance or skewness is large, especially for small sample sizes. In such cases, a Chebyshev UCL with a higher confidence coefficient such as 0.99 may be used, according to Singh, et al.

**Exhibit 12: Steps for Computing UCL Based on the Chebyshev Inequality —  
Nonparametric**

Let  $X_1, X_2, \dots, X_n$  represent the  $n$  randomly sampled concentrations.

STEP 1: Compute the arithmetic mean of the data  $\bar{X} = \frac{1}{n} \sum_{i=1}^n X_i$ .

STEP 2: Compute the sample standard deviation  $s = \sqrt{\frac{1}{n-1} \sum_{i=1}^n (X_i - \bar{X})^2}$ .

STEP 3: Compute the one-sided  $(1-\alpha)$  upper confidence limit on the mean

$$UCL_{1-\alpha} = \bar{X} + \sqrt{\frac{1}{\alpha} - 1} (s / \sqrt{n})$$

**Exhibit 13: An Example Computation of UCL Based on Chebyshev Inequality —  
Nonparametric**

Using the same concentration values given in Exhibit 4 and used in Exhibit 11, the UCL on the mean can also be computed based on the Chebyshev inequality.

STEP 1: The sample mean of the  $n=31$  values is  $\bar{X}=9.59$ .

STEP 2: The sample standard deviation of the values is  $s = 9.094$

STEP 3: The one-sided 95% upper confidence limit on the mean is therefore

$$UCL_{95\%} = 9.59 + 4.3589 \times 9.094 / \sqrt{31} = 16.7$$



## 5.0 OPTIONAL USE OF MAXIMUM OBSERVED CONCENTRATION

Because some of the methods outlined above (particularly the Land method) can produce very high estimates of the UCL, EPA (1992) allows the maximum observed concentration to be used as the exposure point concentration rather than the calculated UCL in cases where the UCL exceeds the maximum concentration.

It is important to note, however, that defaulting to the maximum observed concentration may not be protective when sample sizes are very small because the observed maximum may be smaller than the population mean. Thus, it is important to collect sufficient samples in accordance with the DQOs for a site. The use of the maximum as the default exposure point concentration is reasonable only when the data samples have been collected at random from the exposure unit and the sample size is large.

## 6.0 UCLs AND THE RISK ASSESSMENT

Risk assessors are encouraged to use the most appropriate estimate for the EPC given the available data. The flow chart in Figure 1 provides general guidelines for selecting a UCL calculation method. Exhibit 14 summarizes the methods described in this guidance, including their applicability, advantages and disadvantages. While the methods identified in this guidance may be useful in many situations, they will probably not be appropriate for all hazardous waste sites. Moreover, other methods not specifically described in this guidance may be most appropriate for particular sites. The EPA risk assessor and, potentially, a trained statistician should be involved in the decision of which method(s) to use.

When presenting UCL estimates, the risk assessor should identify:

- how the shape of the underlying distribution was identified (or, if it was not identified, what methods were used in trying to identify it),
- the chosen UCL method,
- reasons that this UCL method is appropriate for the site data, and
- assumptions inherent in the UCL method.

It may also be appropriate to include information such as advantages and disadvantages of the distribution-fitting method, advantages and disadvantages of the UCL method, and how the risk characterization would change if other assumptions were used.

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<b>Exhibit 14</b> <b>Summary of UCL Calculation Methods</b>				
Method	Applicability	Advantages	Disadvantages	Reference
<i>For Normal or Lognormal Distributions</i>				
Student's <i>t</i>	means normally distributed, samples random	simple, robust if <i>n</i> is large	distribution of means must be normal	Gilbert 1987; EPA 1992
Land's <i>H</i>	lognormal data, small variance, large <i>n</i> , samples random	good coverage <sup>1</sup>	sensitive to deviations from lognormality, produces very high values for large variance or small <i>n</i>	Gilbert 1987; EPA 1992
Chebyshev Inequality (MVUE)	skewness and variance small or moderate, samples random	often smaller than Land	may need to resort to higher confidence levels for adequate coverage	Singh et al. 1997
Wong	gamma distribution	second order accuracy <sup>2</sup>	requires numerical solution of an improper integral	Schulz and Griffin 1999; Wong 1993
<i>Nonparametric/Distribution-free Methods</i>				
Central Limit Theorem - Adjusted	large <i>n</i> , samples random	simple, robust	sample size may not be sufficient	Gilbert 1987; Singh et al. 1997
Bootstrap <i>t</i> Resampling	sampling is random and representative	useful when distribution cannot be identified	inadequate coverage for some distributions; computationally intensive	Singh et al. 1997; Efron 1982
Hall's Bootstrap Procedure	sampling is random and representative	useful when distribution cannot be identified; takes bias and skewness into account	inadequate coverage for some distributions; computationally intensive	Hall 1988; Hall 1992; Manly 1997; Schultz and Griffin 1999
Jackknife Procedure	sampling is random and representative	useful when distribution cannot be identified	inadequate coverage for some distributions; computationally intensive	Singh et al. 1997
Chebyshev Inequality	skewness and variance small or moderate, samples random	useful when distribution cannot be identified	inappropriate for small sample sizes when skewness or variance is large	Singh et al. 1997; EPA 2000c
<sup>1</sup> Coverage refers to whether a UCL method performs in accordance with its definition.				
<sup>2</sup> As opposed to maximum likelihood estimation, which offers first order accuracy.				

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## **7.0 PROBABILISTIC RISK ASSESSMENT**

The estimates of the UCL described in this guidance can be used as point estimates for the EPC in deterministic risk assessments. In probabilistic risk assessments, a more complete characterization of the underlying distribution of concentrations may be important as well. Risk assessors should consult *Risk Assessment Guidance for Superfund, Volume 3 - Part A, Process for Conducting a Probabilistic Risk Assessment* (EPA 2001b) for specific guidance with respect to probabilistic risk assessments.

## **8.0 CLEANUP GOALS**

Cleanup goals are commonly derived using the risk estimates established during the risk assessment. Often, a cleanup goal directly proportional to the EPC will be used, based on the relationship between the site risk and the target risk as defined in the National Contingency Plan. In such cases, the attainment of the cleanup goal should be measured with consideration of the method by which the EPC was derived. For more details, see *Surface Soil Cleanup Strategies for Hazardous Waste Sites* (EPA, to be published).

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### Appendix A: Using Bounding Methods to Account for Non-detects

This appendix presents an iterative procedure that can be used to account for non-detects in data when estimating a UCL. It provides a step-by-step approach for computing an upper bound on the UCL using the "Solver" feature in Microsoft® Excel spreadsheets.

STEP 1. Enter all the detected values in a column.

STEP 2. At the bottom of the same column, append as place holders as many copies of the formula

$=\text{RAND}()*DL$

as there were non-detects. In these formulas,  $DL$  should be replaced by the detection limit.

STEP 3. Copy all the cells you have entered in steps 1 and 2 to a second column.

STEP 4. In another cell, enter the formula for the UCL that you wish to use. For instance, to use the 95% UCL based on Student's  $t$ , enter the formula

$=\text{AVERAGE}(\text{range})+\text{TINV}((1-0.95)*2, n-1)*\text{SQRT}(\text{VAR}(\text{range})/n)$

where *range* denotes the array of cell references in the second column you just created and  $n$  denotes the number of measurements (both detected values and non-detects).

STEP 5. From the Excel menu, select Tools / Solver.

STEP 6. In the "Solver Parameters" dialog box, specify the cell in which you entered the UCL formula as the Target Cell.

STEP 7. To find the upper bound of the UCL click on the Max indicator; to find the lower bound of the UCL click on the Min indicator.

STEP 8. Enter references to the cells containing the place holders for the non-detects in the field under the label "By Changing Cells." (Do not click the "Guess" button.)

STEP 9. For each cell that represents a non-detect, add a constraint specifying that the cell is to be greater than or equal to (" $\geq$ ") the detection limit  $DL$ .

STEP 10. Click on the Options button and check the box labeled "Assume Non-Negative."

STEP 11. Then click OK and then the Solver button. The program will automatically locate a local extreme value (i.e., maximum or minimum) for the UCL.

STEP 12. Record this value. You can use the Save Scenario button and Excel's scenario manager to do this.

STEP 13. Again copy all the detected values and randomized place holders for the non-detects from the first column to the same spot in the second column.

STEP 14. Select Tools / Solver and click the Solve button.

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STEP 15. If calculating the upper bound, record the resulting value of the UCL if it is larger than previously computed. If calculating the lower bound, record the resulting value of the UCL if it is smaller than previously computed.

STEP 16. Repeat steps 13 through 15 to search for the global maximum or minimum value for the UCL.



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### Appendix B: Computer Code for Computing a UCL with the Hall's Bootstrap Sampling Method

This appendix presents Pascal code that can be used to compute the bootstrap estimate of a UCL. To use it, place data in the vector  $x$ . Then specify the sample size  $n$ , the vector  $x$  and the alpha-level, and call the procedure bootstrap. When the procedure finishes, the estimated value will be in the variable UCL. To obtain a 95% UCL, let alpha be 0.05. Up to 100 data values and up to 10,000 bootstrap iterations are supported, but these limits may be changed.

```

const
  max = 100;
  bmax = 10000;

type
  index = 1..max;
  bindex = 1..bmax;
  float = extended; {could just be real}
  vector = array[index] of float;
  bvector = array[bindex] of float;

var
  qq : bvector;

function getmean(n : integer; x : vector) : float;
var s : float; i : integer;
begin
  s := 0.0;
  for i := 1 to n do s := s + x[i];
  getmean := s / n;
end;

function getstddev(n:integer; xbar:float; x:vector) : float;
var s : float; i : integer;
begin
  s := 0.0;
  for i := 1 to n do s := s + (x[i] - xbar) * (x[i] - xbar);
  getstddev := sqrt(s / n); {not n-1}
end;

function getskew(n:integer; xbar:float; stddev:float; x:vector) :
float;
var s,s3 : float; i : integer;
begin
  s := 0.0;
  s3 := stddev * stddev * stddev;
  for i:=1 to n do s:=s+(x[i]-xbar)*(x[i]-xbar)*(x[i]-xbar)/s3;
  getskew := s / n;
end;

procedure qsort(var a: bvector; lo,hi: integer);
procedure sort(l,r: integer);
var i,j : integer; x,y: float;
begin
  i:=l; j:=r; x:=a[(l+r) div 2];
  repeat
    while a[i]<x do i:=i+1;
    while x<a[j] do j:=j-1;
    if i<=j then
      begin
        y:=a[i]; a[i]:=a[j]; a[j]:=y;
        i:=i+1; j:=j-1;
      end;
  until i>j;
  sort(l,i);
  sort(j,r);
end;

```

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```

    until i>j;
    if l<j then sort(l,j);
    if i<r then sort(i,r);
end;
begin {qsort}
sort(lo,hi);
end;

procedure bootsample(n : integer; x : vector; var y : vector);
var i,j : integer;
begin
for i := 1 to n do
begin
j := random(n) + 1;
y[i] := x[j];
end;
end;

procedure bootstrap(n:integer; x:vector; alpha:float; var
ucl:float);
{let alpha be 0.05 to compute a 95% UCL}
var
i,b,bb : integer;
xbar, stddev, skew, bxbar, bstddev, bskew, k, w, q, a : float;
bx : vector;
begin
bb := bmax;
for b:=1 to bmax do qq[b] := 0.0;
xbar := getmean(n,x);
stddev := getstddev(n,xbar,x);
skew := getskew(n,xbar,stddev,x);
for b := 1 to bb do
begin
bootsample(n,x,bx);
bxbar := getmean(n,bx);
bstddev := getstddev(n,bxbar,bx);
k := getskew(n,bxbar,bstddev,bx);
w := (bxbar - xbar) / bstddev;
q := w + skew * w*w / 3 + k*k * w*w*w / 27 + k / (6 * n);
qq[b] := q;
end;
qsort(qq,1,bb);
q := qq[round(alpha * bb)];
a := 1 + skew * (q-skew / (6 * n));
if a = 0.0 then w := -3 / skew
else w := (3 / skew) * (exp((1/3) * ln(a)) - 1);
ucl := xbar - w * stddev;
end;

```